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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant: Peter HUDSON et al.

Title: BENZAMIDE DERIVATIVES AS OXYTOCIN
AGONISTS AND VASOPRESSIN
ANTAGONISTS

Appl. No.: 10/541,460

Filing Date: March 6, 2006

Examiner: Unassigned

Art Unit: Unassigned

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT
UNDER 37 CFR §1.56

Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Sir:

Submitted herewith on Form PTO/SB/08 is a listing of documents known to Applicants in order to comply with Applicants' duty of disclosure pursuant to 37 CFR §1.56.

A copy of each non-U.S. patent document and each non-patent document is being submitted to comply with the provisions of 37 CFR §1.97 and §1.98.

The submission of any document herewith, which is not a statutory bar, is not intended as an admission that such document constitutes prior art against the claims of the present application or that such document is considered material to patentability as defined in 37 CFR §1.56(b). Applicants do not waive any rights to take any action which would be appropriate to antedate or otherwise remove as a competent reference any document which is determined to be a *prima facie* art reference against the claims of the present application.

TIMING OF THE DISCLOSURE

The listed documents are being submitted in compliance with 37 CFR §1.97(b), before the mailing date of the first Office Action on the merits.

RELEVANCE OF EACH DOCUMENT

The relevance of the foreign-language documents is described on page 17 of the specification. An English translation of the foreign-language documents is not readily available. However, the absence of such translation does not relieve the PTO from its duty to consider the submitted foreign language documents (37 CFR §1.98 and MPEP §609).

Applicants respectfully request that each listed document be considered by the Examiner and be made of record in the present application and that an initialed copy of Form PTO/SB/08 be returned in accordance with MPEP §609.

Although Applicant believes that no fee is required for this Request, the Commissioner is hereby authorized to charge any additional fees which may be required for this Request to Deposit Account No. 19-0741.

Respectfully submitted,

Date 6 March 2006

By S.A. Bent

FOLEY & LARDNER LLP
Customer Number: 22428
Telephone: (202) 672-5404
Facsimile: (202) 672-5399

Stephen A. Bent
Attorney for Applicant
Registration No. 29,768

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Substitute for form 1449B/PTO

**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**

Date Submitted: March 6, 2006

(use as many sheets as necessary)

Sheet

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of

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Complete if Known

Application Number	10/541,460
Filing Date	March 6, 2006
First Named Inventor	Peter HUDSON
Group Art Unit	Unassigned
Examiner Name	Unassigned

Attorney Docket Number

052209-0138

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number	Kind Code ² (if known)			

U.S. PATENT APPLICATION DOCUMENTS

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FOREIGN PATENT DOCUMENTS

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		Office ³	Number ⁴	Kind Code ⁵ (if known)				

NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.) date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁶
B1		Thomas BOSSMAR et al., "Effects of SR 49059, an orally active V _{1a} vasopressin receptor antagonist, on vasopressin-induced uterine contractions", British Journal of Obstetrics & Gynaecology, April 1997, Vol. 104, pp. 471-477.	
B2		R. BROUARD et al., "Effect of SR49059 an orally active V _{1a} vasopressin receptor antagonist, in the prevention of dysmenorrhoea", British Journal of Obstetrics & Gynaecology , May 2000, Vol. 107, pp. 614-619.	
B3		Venkatesan ARANAPAKAM et al., '4,10-DIHYDRO-5H-THIENO[3,2-c][1]BENZAZEPINE DERIVATIVES AND 9,10-DIHYDRO-4H-THIENO[2,3-C][1]BENZAZEPINE DERIVATIVES AS ORALLY ACTIVE ARGININE VASOPRESSIN RECEPTOR AGONISTS", Bioorganic & Medicinal Chemistry Letters 9 (1999) pp. 1733-1736.	
B4		M. ARTICO et al., "RICERCHE SU SOSTANZE AD ATTIVITA ANTIBLASTICA", Il Fármaco., Ed. Sc. Vol. 24, no. 3, pp. 276-284.	
B5		F. CHIMENTI et al., "RICERCHE SU SOSTANZE AD ATTIVITA ANTIBLASTICA", Il Fármaco, Ed. Sc. Vol. 32, no. 5, pp. 339-347.	

Examiner Signature

Date Considered

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹ Unique citation designation number. ²See attached Kinds of U.S. Patent Documents. ³Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document.⁵Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶Applicant is to place a check mark here if English language Translation is attached.

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	B6	Jiban K. CHAKRABARTI et al., "4-piperazinyl-10H-thieno[2,3-b][1,5]benzodiazepines as Potential Neuroleptics", J. Med. Chem. 1980, 23, pp. 878-884.	
	B7	Jiban K. CHARABARTI et al., "10-Piperazinyl-4H-thieno[3,2-b][1,5]- and -[3,4-b][1,5]benzodiazepines as Potential Neuroleptics", J. Med. Chem. 1980, 23, pp. 884-889.	
	B8	Jiban K. CHARABARTI et al., 'Synthesis and Pharmacological Evaluation of a Series of 4-Piperazinylpyrazolo[3,4-b]- and -[4,3-b][1,5]benzodiazepines as Potential Anxiolytics", J. Med. Chem. 1989, 32, pp. 2573-2582.	
	B9	Alba CHIMIRRI et al., "ANNELATED 1,5-BENZODIAZEPINES. PART I. THREE, FOUR, AND FIVE MEMBERED RINGS", HETEROCYCLES, VOL. 36, NO. 3, 1993, PP. 601-637.	
	B10	Gary L. GUNEWALD et al., "Effect of Ring Size or an Additional Heteroatom on the Potency and Selectivity of Bicyclic Benzylamine-Type Inhibitors of Phenylethanolamine N-Methyltransferase", J. MED. CHEM. 1996, 39, 3539-3546.	
	B11	Janice M. KLUNDER et al., "Novel Non-Nucleoside Inhibitors of HIV-1 Reverse Transcriptase. 2. Tricyclic Pyridobenzoxazepinones and Dibenzoxazepinones", J. Med. Chem., 1992, 35, pp. 1887-1897.	
	B12	Jean-Françoise F. LIÉGEOIS et al., "Pyridobenzoxazepine and Pyridobenzothiazepine Derivates as Potential Central Nervous System Agents: Synthesis and Neurochemical Study", J. Med. Chem. 1994, 37, pp. 519-525.	
	B13	Timothy O. OLAGBEMIRO et al., "Alkylation and an Unusual Reductive Ring Opening of Some Thieno[3,4-b][1,5]benzoxazepin-10-ones", Department of Chemistry, Bayero University, Kano, Nigeria, VOL 19, NOV-DEC 1982, pp. 1501-1504	
	B14	William B. WRIGHT Jr. et al., "Derivatives of 11-(1-Piperazinyl)-5H-pyrrolo[2,1-c][1,4]benzodiazepine as Central Nervous Systems Agents", J. Med. Chem. 1980, 23, pp. 462-465.	
	B15	Satoru SASATANI et al., "DIISOBUTYLALUMINUM HYDRIDE A NOVEL REAGENT FOR THE REDUCTION OF OXIMES", Tetrahedron Letters, Vol. 24, No. 43, pp. 4711-4712, 1983.	

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